## LOCAL CAUSAL DISCOVERY FOR STRUCTURAL EVIDENCE OF DIRECT DISCRIMINATION

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Efficient graph learning enables causal fairness analysis in complex decision systems.

## DETECTING DIRECT DISCRIMINATION == CAUSAL PARENT DISCOVERY



Fig. 1: The standard fairness model (SFM) with protected attribute X, outcome Y, confounders C, and mediators M (bidirected edges denote latent confounding) [1]. We can project the true causal DAG onto the SFM to facilitate fairness analysis. This work identifies direct mechanisms of unfairness in a data-driven way by first discovering  $M \cup C$ .



# LD3: CAUSAL PARENT DISCOVERY FOR FAIRNESS ANALYSIS.

no observed descendants and no unobserved parents (other latent variables are permitted).



Fig. 2: The nodes of any DAG can be uniquely partitioned into 8 disjoint subsets defined by the paths shared with a given pair  $\{X, Y\}$  [2]. This applies to DAGs of any size; triple DAGs are for illustration only. Partition  $\mathbf{Z}_1$  generalizes the confounder,  $\mathbf{Z}_2$  the collider,  $\mathbf{Z}_3$  the mediator, etc.

- FAIRNESS CRITERIA. LD3 results directly evaluate the SDC and can be used as a valid adjustment set for the WCDE:

Definition 1 (Structural direct criterion (SDC), Plečko and Bareinboim 2024). A structural causal model is fair w.r.t. direct discrimination if and only if the following evaluates to 0:

> $SDC = \mathbf{1}(X \in parents(Y)).$ (1)

## **RESULTS**

- FASTER. LD3 ran 46–5870× faster than baselines on real-world data.
- ENABLES EFFECT ESTIMATION. LD3 returns a valid adjustment set for the WCDE under a new graphical criterion.



Fig. 3: Baseline results for parent discovery on the SANGIOVESE benchmark (bnlearn). Independence test count (Tests) is reported for constraintbased methods. Time is in seconds. Shaded regions denote 95% confidence intervals over ten replicates.

# CASE STUDY: LIVER TRANSPLANT ALLOCATION

**Fairness query:** Are sex-based disparities due to direct discrimination?  $\Rightarrow$  **Graphical query:** Is patient sex (*S*) a parent of liver allocation (*L*)?



#### **REFERENCES**

• APPROACH. We introduce LD3, a constraint-based discovery method that leverages the causal partition taxonomy proposed in [2] to label variables by their causal relation to the protected attribute X and outcome Y (Fig. 2), rather than learning the full graph. We assume that Y has

#### • COMPLEXITY. LD3 discovers parents(Y) ( $\in \mathbb{Z}_1 \cup \mathbb{Z}_3 \cup \mathbb{Z}_4$ ) in a linear number of conditional independence tests w.r.t. variable set size.

**Definition 2** (Weighted controlled direct effect (WCDE), Pearl 2000). Let  $\mathbf{M}' \subseteq \mathbf{M}$ denote mediators that are parents of Y. WCDE is a qualitative indicator of direct discrimination, as it is nonzero if and only if  $X \in parents(Y)$  (i.e., SDC = 1):

WCDE = 
$$\sum_{\mathbf{m}'} \left( \mathbb{E}[Y \mid do(x, \mathbf{m}')] - \mathbb{E}[Y \mid do(x^*, \mathbf{m}')] \right) P(\mathbf{m}').$$
(2)

MORE PLAUSIBLE RESULTS. Parent sets predicted from real-world data aligned with expert knowledge better than baselines.







Fig. 4: Predicted parent sets for OPTN STAR datasets ('17-'19, '20-'22). Known parents of L are in yellow. Exposure = patient sex (S; red), outcome = receiving a liver (L; blue). AE = active exception case; BT = blood type; DX = diagnosis; ED = education; ET = ethnicity; EX = exception type; IA = initial age; IM = initial MELD; PM = payment method; RE = region; WE = weight. For all methods, SDC = 1 and WCDE p-value = 0.000.

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